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1		AN	ACT relating to drug disposal.
2	Be it enacted by the General Assembly of the Commonwealth of Kentucky:		
3		→ S	ECTION 1. A NEW SECTION OF KRS CHAPTER 205 IS CREATED TO
4	REA	AD AS	S FOLLOWS:
5	<u>(1)</u>	As ı	used in this section, "drug disposal method" means a nontoxic composition
6		<u>for</u>	the sequestration, deactivation, destruction, and disposal of any unused,
7		unw	anted, or expired prescription.
8	<u>(2)</u>	Upo	n the request of a pharmacist or practitioner as defined in KRS 218A.010, the
9		<u>Ken</u>	tucky Medicaid program shall provide payment to the pharmacist or
10		<u>prac</u>	titioner for the cost of a drug disposal method that the pharmacist or
11		<u>prac</u>	titioner distributed to a Medicaid recipient upon dispensing a controlled
12		subs	stance that contains any salt, compound, derivative, or preparation of an
13		<u>opio</u>	id, benzodiazepine, a barbiturate, codeine, or an amphetamine to the
14		Mea	licaid recipient, in accordance with Section 2 of this Act.
15	<u>(3)</u>	The	amount of the payment shall be the cost to the pharmacist or practitioner of
16		the c	drug disposal method.
17	<u>(4)</u>	The	pharmacist or practitioner shall only receive payment for the cost of the drug
18		<u>disp</u>	osal method if it was provided at no cost to the Medicaid recipient.
19		→ S	ection 2. KRS 218A.170 is amended to read as follows:
20	(1)	A d	uly licensed manufacturer, distributor, or wholesaler may sell or distribute
21		cont	rolled substances, other than samples, to any of the following persons:
22		(a)	To a manufacturer, wholesaler, or pharmacy;
23		(b)	To a practitioner;
24		(c)	To the administrator in charge of a hospital, but only for use by or in that
25			hospital;
26		(d)	To a person in charge of a laboratory, but only for use in that laboratory for
27			scientific and medical research purposes;

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1 (e) To a person registered pursuant to the federal controlled substances laws.

- 2 (2) A pharmacist may sell or distribute a controlled substance:
- 3 (a) Pursuant to a prescription that conforms to the requirements of this chapter; or
- 4 (b) To a person registered pursuant to the federal controlled substances laws.
- 5 (3) A pharmacist who is licensed under KRS Chapter 315 or a pharmacist's designee
- 6 shall inform persons who receive a prescription for a controlled substance that
- 7 contains any salt, compound, derivative, or preparation of an opioid,
- 8 benzodiazepine, a barbiturate, codeine, or an amphetamine, about the importance of
- 9 proper and safe disposal of unused, unwanted, or expired prescription drugs by one
- of the following methods:
- 11 (a) Verbally;
- 12 (b) In writing; or
- 13 (c) Posted signage.
- 14 (4) Upon dispensing of any prescription that contains any salt, compound, derivative,
- or preparation of an opioid, benzodiazepine, a barbiturate, codeine, or an
- amphetamine, a pharmacist who is licensed under KRS Chapter 315 or a
- pharmacist's designee may:
- 18 (a) Make available for purchase, or at no charge distribute, a nontoxic
- composition for the sequestration, deactivation, destruction, and disposal of
- any unused, unwanted, or expired prescription; or
- 21 (b) Provide an on-site, safe, and secure medicine disposal receptacle or kiosk for
- 22 the safe disposal of any unused, unwanted, or expired prescription.
- 23 (5) A manufacturer or distributor of nontoxic compositions for the sequestration,
- deactivation, or destruction and disposal of controlled substances is strongly
- encouraged to enter into a consignment-reimbursement contract with a pharmacy in
- order for a pharmacy to expand its inventory of the nontoxic compositions.
- 27 (6) A practitioner may:

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1		(a)	Administer, dispense, or prescribe a controlled substance only for a legitimate	
2			medical purpose and in the course of professional practice; or	
3		(b)	Distribute a controlled substance to a person registered pursuant to the federal	
4			controlled substance laws.	
5	(7)	A p	ractitioner who dispenses a controlled substance that contains any salt,	
6		compound, derivative, or preparation of an opioid, benzodiazepine, a barbiturate,		
7		code	ine, or an amphetamine shall:	
8		(a)	Inform all persons who receive a prescription for a controlled substance about	
9			the importance of proper and safe disposal of unused, unwanted, or expired	
10			prescription drugs; and	
11		(b)	Make available for purchase, or at no cost distribute, a nontoxic composition	
12			for the sequestration, deactivation, or destruction and disposal of unused,	
13			unwanted, or expired controlled substances.	
14	(8)	All	sales and distributions shall be in accordance with KRS 218A.200 and the	
15		feder	ral controlled substances laws, including the requirements governing the use of	
16		orde	r forms.	
17	(9)	Possession of or control of controlled substances obtained as authorized by this		
18		secti	on shall be lawful if in the regular course of business, occupation, profession,	
19		empl	loyment, or duty of the possessor.	
20	(10)	Subs	sections (3), (4), (7), and (12) of this section shall not apply to veterinarians.	
21	(11)	The	Kentucky Medicaid program shall [not be required to] provide payment for the	
22		prov	isions established in subsections (4) and (7) of this section <i>in accordance with</i>	
23		Secti	ion 1 of this Act.	
24	(12)	Any	person who violates subsection (3) or (7) of this section shall be subject to a	
25		fine	of twenty-five dollars (\$25) for the first violation, a fine of one hundred dollars	
26		(\$10	0) for the second violation, and a fine of two hundred dollars (\$200) for each	
27		subs	equent violation.	

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→ Section 3. If the Department for Medicaid Services or the Cabinet for Health and Family Services determines that a state plan amendment, waiver, or any other form of authorization or approval from a federal agency is necessary prior to the implementation of Sections 1 and 2 of this Act for any reason, including the loss of federal funds, the department shall, within 90 days after the effective date of this Act, request the state plan amendment, waiver, authorization, or approval, and may only delay full implementation of those provisions for which a state plan amendment, waiver, authorization, or approval was deemed necessary until the state plan amendment, waiver, authorization, or approval is granted.